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	Page 1	
1	IN THE UNITED STATES DISTRICT COURT	09:23:02
	SOUTHERN DISTRICT OF CALIFORNIA	
2		09:23:02
3		09:23:02
	X	
4	)	09:23:02
	GEN-PROBE INCORPORATED, )	
5	) NO.99cv2668 H (AJB)	09:23:02
	Plaintiff, )	
6	vs.	09:23:02
	)	
7	VYSIS, INC.,	09:23:02
	)	
8	Defendant. )	09:23:02
	)	
9	X	09:23:02
10		09:23:02
	CONFIDENTIAL	09:23:02
11		09:23:02
12	Videotaped Deposition of	09:23:02
13	JONATHON MICHAEL LAWRIE, Ph.D.	09:23:02
14	Durham, North Carolina	09:23:02
15 16	Thursday, February 15, 2001	09:23:02
17		09:23:02
18	Reported by:	09:23:02
10	Sydney C. Silva, Registered Professional Reporter	
19	File No:	09:23:02
20		
21		
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23	Ex. 9 Pg. 45	
24		
	Ex. 7 Pg. 45	



Reported by: Sydney C. Silva Spherion Deposition Services (704) 333-9889 Fax (704) 372-4593

## Gen-Probe, Incorporated Vysis, Inc. Jonathon Michael Lawrie, PhD

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1	A. No.	14:34:16
2	Q. Do you know whether there's any reference	14:34:17
3	in the patent to a combination of target capture	14:34:19
4	with a target-specific method of application of	14:34:25
5	amplification?	14:34:26
6	A. This patent here?	14:34:27
7	Q. Yes.	14:34:29
8	A. I haven't read it completely, just the	14:34:29
9	pieces you have shown me.	14:34:32
10	Q. When the patent application was filed,	14:34:34
11	did you have any impression about whether the	14:34:36
12	greatest degree of specificity sensitivity might be	14:34:38
13	obtained by combining target capture with a	14:34:42
14	target-specific method of amplification?	14:34:47
15	A. I don't remember.	14:34:56
16	Q. Does that stand to reason at all?	14:34:56
17	A. I don't think so. I don't know what the	14:34:58
18	thought process would have been back then.	14:35:00
19	Q. Can you recall any reason that a	14:35:03
20	reference to PCR might have been intentionally	14:35:05
21	omitted from the patent application?	14:35:08
22	A. Yes.	14:35:15
23	Q. And what reason was that? Let me, let me	14:35:15
24	start over. Ex. 9 Pg. 46	14:35:23





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1	Was a reference to PCR intentionally	14:35:24
2	omitted from the patent to the best of your	14:35:27
3	understanding?	14:35:29
4	A. I don't know.	14:35:30
5	Q. Were there discussions about whether or	14:35:31
6	not to include a reference to PCR in the patent?	14:35:32
7	A. I can't remember.	14:35:36
8	Q. So at Amoco you had a thought about	14:35:47
9	combining target capture with PCR, is that right?	14:35:51
10	A. Yes.	14:35:54
11	Q. Gene-Trak then did work in an effort to	14:35:55
12	combine target capture with PCR, is that right?	14:35:58
13	A. From seeing this here, yes.	14:36:03
14	Q. Do you have a recollection of that?	14:36:05
15	A. No.	14:36:07
16	Q. If there's no reference in the patent to	14:36:07
17	combining target capture with PCR, do you have any	14:36:09
18	explanation as to why it is not there?	14:36:13
19	A. I believe that it was a separate the	14:36:15
20	thought behind this was coming up with new methods	14:36:17
21	of amplification, not old ones.	14:36:19
22	Q. And you would, for the purposes of what	14:36:31
23	you just said, you classify PCR as an old method of	14:36:32
24	amplification?	14:36:36
	Ex. 9 Pg. 47	



14:37:38

14:37:40

14:37:41

14:37:49

14:37:50

## Gen-Probe, Incorporated Vysis, Inc. Jonathon Michael Lawrie, PhD

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A. PCR itself was described in the patent,	14:36:37
yes, issued patent.	14:36:40
Q. And your understanding of the 338 patent	14:36:41
was that it was directed to other methods of	14:36:44
amplification?	14:36:47
A. The, it was, it was directed to the	14:36:48
methods disclosed by, you know, the methods	14:36:54
separate from PCR.	14:36:59
Q. Those being the methods, for example, as	14:37:07
the methods set forth in Example 6 and 7?	14:37:10
A. Yes.	14:37:14
Q. Is it your understanding that the 338	14:37:20
patent then doesn't encompass the combination of	14:37:22
target capture and PCR?	14:37:28
MR. BANKS: Object to the form.	14:37:30
A. I couldn't say.	14:37:31
Q. I'm sorry?	14:37:32
A. I couldn't say.	14:37:32
Q. Was it your intention that it encompass	14:37:33
	A. PCR itself was described in the patent, yes, issued patent. Q. And your understanding of the 338 patent was that it was directed to other methods of amplification? A. The, it was, it was directed to the methods disclosed by, you know, the methods separate from PCR. Q. Those being the methods, for example, as the methods set forth in Example 6 and 7? A. Yes. Q. Is it your understanding that the 338 patent then doesn't encompass the combination of target capture and PCR? MR. BANKS: Object to the form. A. I couldn't say. Q. I'm sorry? A. I couldn't say.

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Q. However, your recollection of why -- of if there's no -- your explanation of why there

Ex. 9 Pg. 48

the combination of target capture and PCR?

I don't know.

intention was in regards to PCR.



I can't remember what the

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1	might not be a reference to PCR in the patent is	14:37:53
2	that the patent wasn't intended to cover old	14:37:56
3	methods of amplification such as PCR; is that	14:38:03
4	right?	14:38:06
5	A. The patent was intended to cover the	14:38:07
6	discoveries by myself, Halbert and King that there	14:38:09
7	should be in some, you know, disclosure back at	14:38:15
8	Amoco. That's what the patent was about.	14:38:16
9	why PCR was left out I can just	14:38:22
10	speculate. It wasn't what we came with, it was in	14:38:26
11	the previous, it was a previous older method.	14:38:30
12	Q. You were looking for other things?	14:38:33
13	A. Yeah.	14:38:36
14	MR. BOWEN: Let's assume that the patent	14:39:04
15	application for the 330 patent was filed on	14:39:06
16	December 21, 1987. Can we stipulate to that?	14:39:10
17	MR. BANKS: For which patent?	14:39:16
18	MR. BOWEN: The 330.	14:39:18
19	MR. BANKS: The 330? Moving to a	14:39:20
20	different one now?	14:39:21
21	MR. BOWEN: I'm confused this late in the	14:39:22
22	day, huh? The first application that claimed	14:39:25
23	the combination of target capture and	14:39:27
24	amplification.	14:39:32
	Ex. 9 Pg. 49	

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[	D . 221	
1	Page 231 Example 5 is a linear method?	16:21:41
2	A. Let's see.	16:21:44
3	Yes, it is linear.	16:22:29
4	Q. So Example 5 discloses a linear	16:22:31
5	nonspecific method of amplification?	16:22:34
6	A. Yes.	16:22:37
7	Q. So recapping the examples, Examples 1	16:22:38
8	through 3 disclose capture methods without	16:22:43
9	amplification?	16:22:46
10	A. Yes.	16:22:48
11	Q. And Example 4 discloses linear	16:22:49
12	nonspecific amplification?	16:22:53
13	A. Yes.	16:22:54
14	Q. Example 5 discloses linear nonspecific	16:22:55
15	amplification?	16:22:59
16	A. Yes.	16:23:00
17	Q. Example 6 seeks to describe nonspecific	16:23:02
18	exponential amplification?	16:23:10
19	A. Let's see. Yes.	16:23:13
20	Q. And Example 7 describes seeks to	16:23:18
21	describe nonspecific exponential amplification?	16:23:22
22	A. Yes.	16:23:28
23	Q. Looking back at Column 30, specifically	16:23:44
24	at Lines 30 through 40, which I think is two	16:23:48
	Ex. 9 Pg. 50	

